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- (v) Provided the requirements of paragraph (c)(2)(iv) of this section are met, use the gas divider with the gas selected in paragraph (c)(2)(iii) of this section and determine the remainder of the calibration points. Fit a calibration curve per §§ 90.316, 90.317, 90.318, and 90.320 of this chapter for the entire analyzer range.
- (d) Emission measurement accuracy—continuous sampling. Analyzers used for continuous analysis must be operated such that the measured concentration falls between 15 and 100 percent of fullscale chart deflection. Exceptions to these limits are:
- (1) The analyzer's response may be less than 15 percent or more than 100 percent of full scale if automatic range change circuitry is used and the limits for range changes are between 15 and 100 percent of full-scale chart deflection:
- (2) The analyzer's response may be less than 15 percent of full scale if:
- (i) The alternative in paragraph (c)(2) of this section is used to ensure that the accuracy of the calibration curve is maintained below 15 percent; or
- (ii) The full-scale value of the range is 155 ppm (C) or less; or
- (iii) The emissions from the engine are erratic and the integrated chart deflection value for the cycle is greater than 15 percent of full scale; or
- (iv) The contribution of all data read below the 15 percent level is less than 10 percent by mass of the final test results

## § 90.315 Analyzer initial calibration.

- (a) Warming-up time. The warming-up time should be according to the recommendations of the manufacturer. If not specified, a minimum of two hours should be allowed for warming up the analyzers.
- (b) NDIR, FID, and HFID analyzer. Tune and maintain the NDIR analyzer per the instrument manufacturer recommendations or specifications or using good engineering practice. The combustion flame of the FID or HFID analyzer must be optimized in order to meet the specifications in \$90.316(b).
- (c) Zero setting and calibration. Using purified synthetic air (or nitrogen), set the CO,  $CO_2$ ,  $NO_X$ . and HC analyzers at zero. Connect the appropriate cali-

brating gases to the analyzers and record the values. Use the same gas flow rates and pressure as when sampling exhaust.

(d) Rechecking of zero setting. Recheck the zero setting and, if necessary, repeat the procedure described in paragraph (c) of this section.

## § 90.316 Hydrocarbon analyzer calibration.

- (a) Calibrate the FID and HFID hydrocarbon analyzer as described in this section. Operate the HFID to a set point  $\pm 5.5$  °C between 185 and 197 °C.
- (b) Initial and periodic optimization of detector response. Prior to initial use and at least annually thereafter, adjust the FID and HFID hydrocarbon analyzer for optimum hydrocarbon response as specified in this paragraph. Alternative methods yielding equivalent results may be used, if approved in advance by the Administrator.
- (1) Follow good engineering practices for initial instrument start-up and basic operating adjustment using the appropriate fuel (see §90.312) and purified synthetic air or zero-grade nitrogen.
- (2) Use of one of the following procedures is required for FID or HFID optimization:
- (i) The procedure outlined in Society of Automotive Engineers (SAE) paper No. 770141, "Optimization of a Flame Ionization Detector for Determination of Hydrocarbon in Diluted Automotive Exhausts;" author, Glenn D. Reschke. This procedure has been incorporated by reference. See §90.7.
- (ii) The HFID optimization procedures outlined in 40 CFR part 1065, subpart D.
- (iii) Alternative procedures may be used if approved in advance by the Administrator.
- (3) After the optimum flow rates have been determined, record them for future reference.
- (c) Initial and periodic calibration. Prior to initial use and monthly thereafter, or within one month prior to the certification test, the FID or HFID hydrocarbon analyzer must be calibrated on all normally used instrument ranges using the steps in this paragraph. Use the same flow rate and pressures as when analyzing samples. Introduce